

Response Under 37 CFR §1.116

Expedited Procedure

Examining Group 1652

Application No. 10/538,423

Paper Dated: February 9, 2011

In Reply to USPTO Correspondence of November 9, 2010

Attorney Docket No. 4544-051674

AMENDMENTS TO THE CLAIMS

This listing of claims will replace all prior versions and listings of claims in the application.

Listing of Claims

Claim 1 (Previously Presented): An isolated nucleic acid molecule for a salt-tolerant L-myo-inositol 1-phosphate synthase from *Porteresia coarctata* PcINO1 comprising the nucleic acid sequence of SEQ ID NO: 1 or a nucleic sequence encoding a protein comprising SEQ ID NO: 3.

Claim 2 (Cancelled).

Claim 3 (Previously Presented): A process of obtaining cDNA, encoding a salt-tolerant L-myo-inositol 1-phosphate synthase comprising:

(i) isolation of a full-length cDNA for the L-myo-inositol 1-phosphate synthase gene from the leaf of *Porteresia coarctata* by reverse transcription followed by polymerase chain reaction; and

(ii) sequencing of the isolated L-myo-inositol 1-phosphate synthase gene, wherein the sequenced synthase from *Porteresia coarctata* PcINO1 is encoded by the nucleotide sequence-SEQ ID NO: 1 and has a deduced amino acid sequence SEQ ID NO: 3.

Claim 4 (Previously Presented): The process as claimed in claim 3, wherein the isolated full-length cDNA of PcINO1 is cloned into a suitable bacterial expression vector pET 20B(+) to produce an expression plasmid.

Claim 5 (Currently Amended): The process as claimed in claim 4, wherein said plasmid is introduced into the host strain E. coli BL-21 (DE 3) thereby forming a transformed host strain and wherein the transformed host strain is cultured by culturing the transformed host strain to express the PcINO1 gene product.

Response Under 37 CFR §1.116

Expedited Procedure

Examining Group 1652

Application No. 10/538,423

Paper Dated: February 9, 2011

In Reply to USPTO Correspondence of November 9, 2010

Attorney Docket No. 4544-051674

Claim 6 (Currently Amended): The process as claimed in claim 5, wherein the expressed PINO1-PcINO1proteins are solubilized in a solubilization buffer containing 8M Urea, 0.5 M NaCl, 20 mM Tris-HCl, pH 7.5,10 mM ME and 2 mM PMSF.

Claim 7 (Previously Presented): A plasmid comprising the isolated nucleic acid molecule of claim 1.

Claim 8 (Previously Presented): A bacteria comprising the isolated nucleic acid molecule of claim 1.